SO-Sequence Ontology Update: Results of Sequence Ontology Workshop on Non-canonical features

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> BRC4 Dec 2006 Jessie Kissinger Jkissing@uga.edu

Participants

- Michael Ashburner
- Suzi Lewis
- Karen Eilbeck
- Ariane Toussaint
- Anne Summers
- Jessie Kissinger











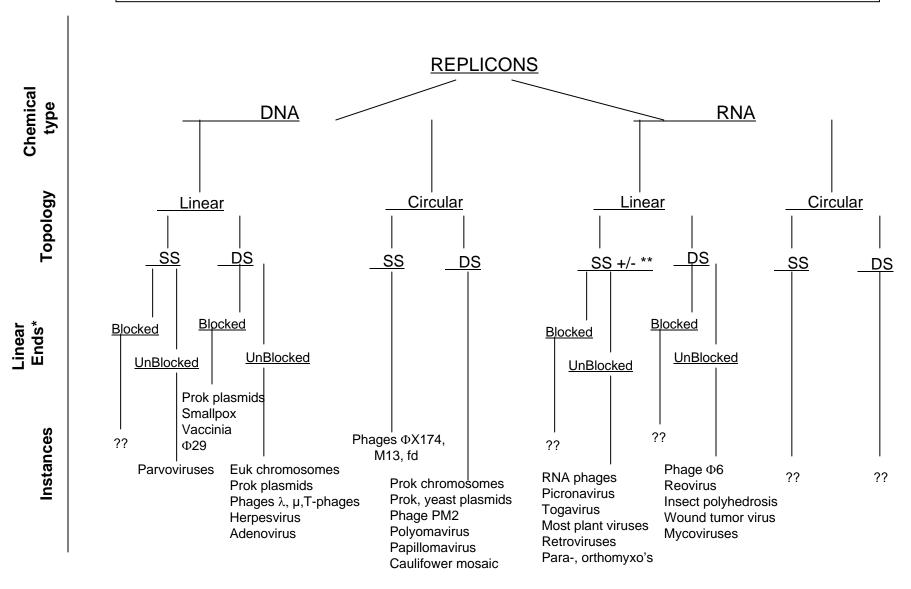


chromocomo variation

| SO home | Releases | CVS | Mailing List | Trackers |

chromosome SO:0000340 Search for terms in SO. Definition: Structural unit composed of long DNA molecule. Type a word to search for in SO. The autocomplete function DBxref:external reference returns a list of matches. [http:///biotech.icmb.utexas.edu/search/dict-search.mhtml] Synonyms: chromosome Aspect: located_sequence_feature Parent relationships: Child relationships: chromosome is a region chromosome region part of chromosome locate term chromosome_arm SO:0000105 **Definition:** A region of the chromosome between the centromere and the telomere. **Browse SO** Human chromosomes have two arms, the p arm (short) and the q arm (long) which are separated from each other by the centromere. miSO provides a graphical means to browse the terms in the DBxref:external reference [http:///www.exactsciences.com/cic/glossary/_index.htm] Sequence Ontology. The view is organised via a parent/child approach. The terms are all linked to the SO term tables web Synonyms: page which displays more details such as the relationship Aspect: located_sequence_feature types, the definition, the synonyms and the cross references. Parent relationships: Child relationships: chromosome arm is a chromosome region Sequence Ontology in the consequences of mutation

A HIERARCHY OF REPLICONS BASED ON CHEMISTRY AND TOPOLOGY



^{*}Ends of linear replicons can either be free (unblocked) 5'phosphates and 3' hydroxyls or blocked in short hairpins (in SS) or covalently crosslinked (DS).

^{**} SS RNA replicons can either be directly translatable mRNA's (+) or complementary to mRNA (-), needing to be copied before used as mRNA.

Basic and Mosaic MGE's

Agents of HGT, MGE (pure prototypes)	Plasmids (RP4, RK2, R100, ColE1)	Phages (lambda)	Transposons (Tn5, Tn10, Tn501)
Mosaic MGE's	P1 A temperate (lysogenic) phage that is a plasmid	Mu A temperate (lysogenic) phage that that replicates by transposition	Conjugative Transposon A transposon that conjugates to other cells, e.g. Tn916
	Genomic Islands have conjugative, phage-derived, and transposon-derived components all located together on the main chromsome. Pathogenicity islands (PAIs) have virulence genes. Also called ICE, integrative conjugative element (20KB - >100s KB)		

MGE's have 3 defining processes

		1. Self-Replication	2. Transfer	3. Site-specific Recombination
Genes	Trans-acting: Enzymes and NA binding proteins	Replicases, NA polymerases, rolling circle replication, phage 'host-takeover' functions	DNA "pump", a Type IV secretion system (T4SS). Phage packaging proteins	Transposases, integrases, excisases, resolvases
	Cis-acting: NA interaction or cleavage sites	oriV	oriT, phage packaging sites, cos site	Gene cassettes, Att sites, Tnp IR's, Res sites

MGE's also carry "baggage" loci: functions that are valuable but not intrinsic to being a gene transfer agent.

Examples of "baggage" genes		1. Pathogenicity	2. Metabolism	3. Resistance
Genes	Trans-acting: Enzymes, transport proteins and NA binding proteins	Toxins , invasion proteins, colonization factors	Catabolic pathways for xenobiotic compounds. Nitrogen fixation, photosynthesis	Antibiotic resistance, toxic metal resistance.
	Cis-acting: NA interaction sites	Relevant operators, promoters, etc.	Relevant operators, promoters, etc	Relevant promoters, operators, etc

Some "baggage" loci reside within transposons that are carried by plasmids or phages.

Ontology Needs & Benefits for Mobile Genetic Elements

• Missing gene ontologies for intrinsic MGE functions, e.g.:

- DNA pump (variant of general secretion systems)
- Recombinases (transposases, resolvases, integrases)
- Phage structural/packaging genes
 - Tail fibers
 - Capsid proteins (self-assemble into phage protein coat)

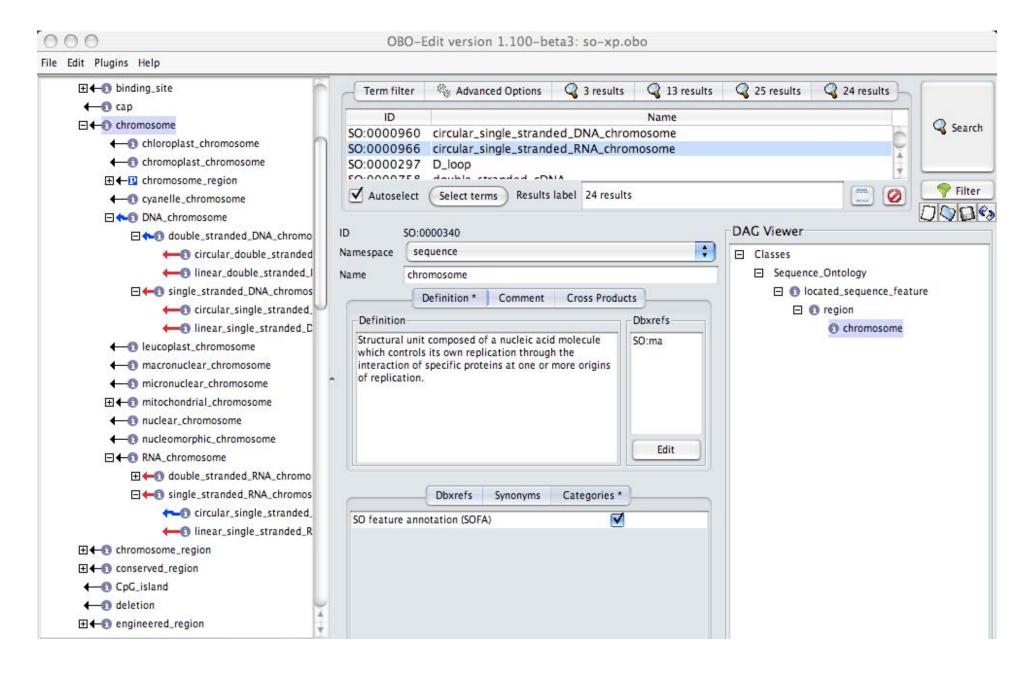
Missing <u>sequence</u> ontologies for intrinsic MGE functions, e.g.:

- Transfer origins, oriT and unique replication origins, oriV
- Res sites for transposition
- Att sites for integration
- Phage packaging sites

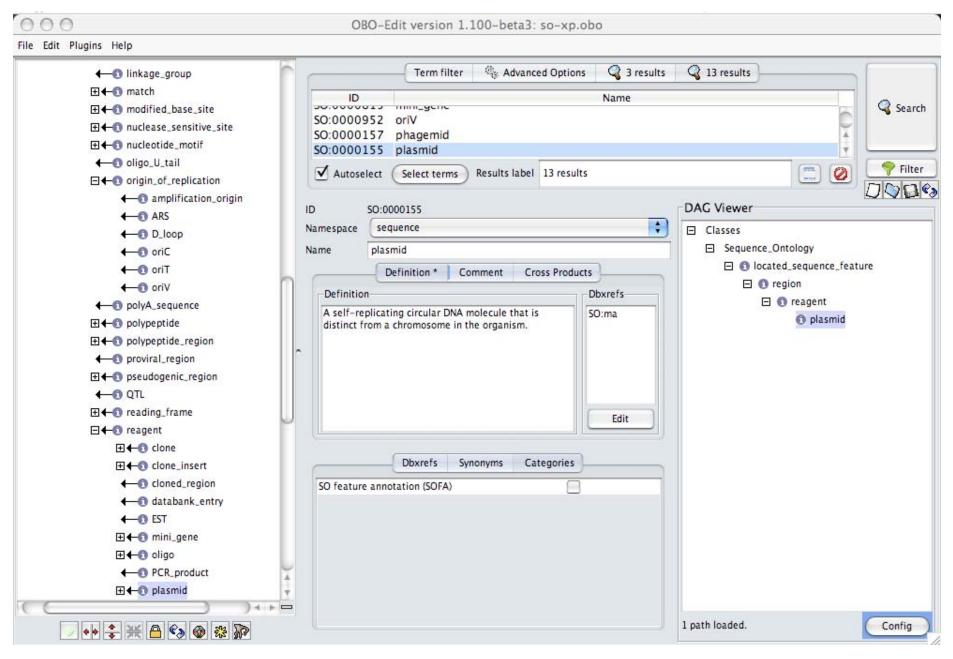
Benefits of improved MGE ontology for the community:

- Literature searching
- Find novel relationships, insights
- Facilitate expert annotation A.I. makes it easier and faster to incorporate their expertise

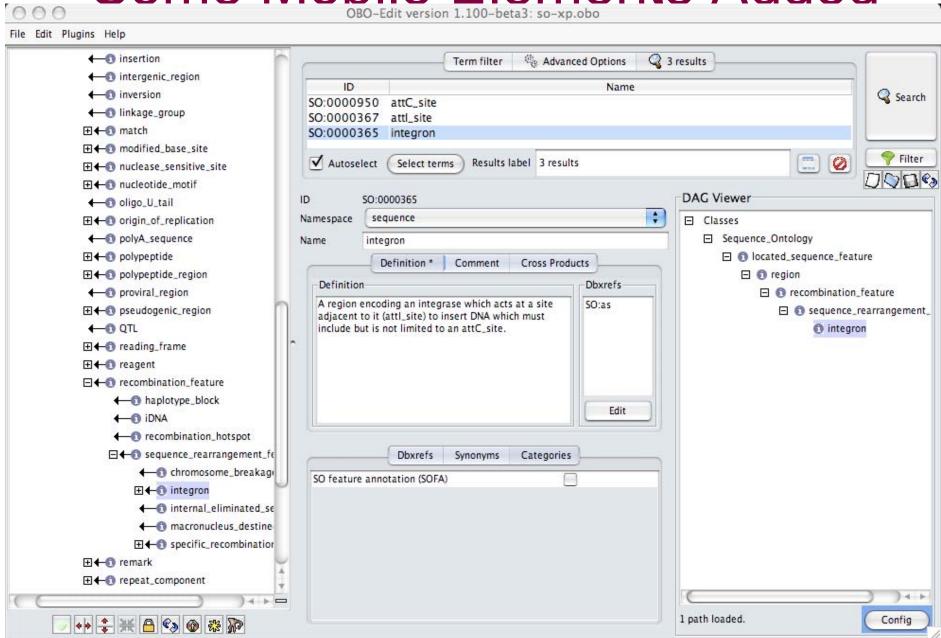
New Chromosome Definintion



Plasmids, not just reagents any more



Some Mobile Elements Added



How to get new terms

- Make a few ppt slides
 - Define the biological problem, pictures help
 - Define the existing terms and their shortcomings
 - Propose new terms and their definitions
- Send the above to Karen Eilbeck, kareneilbeck@mac.com
- Get OBO-Edit from Sourceforge.net download current SO from:
 - http://www.sequenceontology.org/so.shtml

Our current use of SO

- Our usage of available terms is sparse and disparate between BRCs
- BRCs use ~14 terms in our GFF files:

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(contig, mRNA, tRNA, rRNA, gene, CDS, gene, five_prime_utr, three_prime_utr, region, exon, repeat_region, mature_peptide, ribosome_entry_site)
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- GenBank has 50+ terms
- SO has 200+ terms (SOFA has ~100)
- Should we decide on standards and depth? Just because we can annotate it, should we?
- What do our communities need?
- View Sample GFF -SOFA term variation